Art Unit: 1775

EXAMINER'S AMENDMENT

1. An examiner's amendment to the record appears below. Should the changes and/or additions be unacceptable to applicant, an amendment may be filed as provided by 37 CFR 1.312. To ensure consideration of such an amendment, it MUST be submitted no later than the payment of the issue fee.

Authorization for this examiner's amendment was given in a telephone interview with Richard D. Weir on September 27, 2004.

The application has been amended as follows:

IN THE CLAIMS

- 1. (Currently amended) A [An] method for making an electrical-energy-storage unit comprising [of] components fabricated by the method steps as follow;
 - a) preparing a wet-chemical-prepared calcined composition-modified barium titanate powder derived from a solution of precursors: Ba(NO₃)₂, Ca(NO₃)₂•4H₂O, Nd(NO₃)₃•6H₂O, Y(NO₃)₃•4H₂O, Mn(CH₃COO)₂•4H₂O, ZrO(NO₃)₂, and [CH₃CH(O-)COONH₄]₂Ti(OH)₂ in deionized water heated to 80° C, and a separate solution of (CH₃)₄NOH made in deionized water and heated to 80°-85° C, then [by] mixing the solutions by pumping the heated ingredient streams simultaneously through a coaxial fluid mixer producing coprecipitated powder, then collecting the coprecipitated powder [is collected] in a drown-out vessel and [refluxed] refluxing at a temperature of 90°-95° C for 12 hours, then [filtered] filtering, washing with deionized-water [washed], [dryed] drying, and then [calcined] calcining 1050° C in air; and

Art Unit: 1775

- b) <u>fabricating an aluminum oxide (Al₂O₃) coating of 100 Å thickness</u> onto the wet-chemical-prepared calcined composition-modified barium titanate powder, [an aluminum oxide (Al₂O₃) coating of 100 Å thickness was fabricated] with the use of aluminum nitrate nonahydrate precursor applied by wet chemical means, then [calcined] <u>calcining</u> at 1050° C, resulting in a single-coated calcined composition-modified barium titanate powder; and
- c) <u>fabricating</u> onto the alumina-coated composition-modified barium titanate powder, a second uniform coating of 100 Å of calcium magnesium aluminosilicate glass derived [and fabricated] from alcohol-soluble precursors: calcium methoxide or calcium isopropoxide, magnesium methoxide or magnesium ethoxide, aluminum ethoxide or aluminum isopropoxide or aluminum isopropoxide, and tetraethyl orthosilicate are applied by wet chemical means which upon calcining at 500° C results in a double-coated composition-modified barium titanate powder; and
- d) blending, this double-coated composition-modified barium titanate powder
 with a screen-printing ink containing appropriate plastic resins, surfactants,
 lubricants, and solvents to provide a suitable rheology for screen printing; and
- e) screen-printing [printed] into interleaved multilayers of alternating offset nickel electrode layers 12 and double-coated calcined composition-modified barium titanate high-relative-permittivity layers 11 with the use of screening inks having the proper rheology for each of the layers; and

Art Unit: 1775

Page 4

- f) [dry] drying and [cut] cutting the screen-printed multilayer components 15 into a specified rectangular area; and
- g) [binder-burnout and sinter] sintering the screen-printed multilayer components
 15, first at a temperature of 350° C for a specified length of time, then at 850° C for a specified length of time, to form closed-pore porous ceramic bodies;
 and
- h) hot isostatically [press] <u>pressing</u> the closed-pore porous ceramic bodies, at a temperature of 700° C with a specified pressure, into a void-free condition; and
- i) [grind] grinding and [polish] [polishing] each side of the component to expose the alternating offset interleaved nickel electrodes 12; and
- j) connecting nickel side bars 14 [are connected] to each side of the components 15, that have the interleaved and alternating offset nickel electrodes 12 exposed, by applying nickel ink with the proper rheology to each side and clamping the combinations together; and
- k) heating the components and side nickel bar combination 14-15 [are then heated at the proper temperature,] 800° C, and time duration of 20 minutes to bond them together; and
- l) wave [solder] soldering each side of the conducting bars; and
- m) <u>assembling</u> the components 15 with the connected nickel side bars 14 [are then assembled] into the first array, utilizing unique tooling and solder-bump technology; and
- n) assembling the first arrays [are then assembled] into the second array; and

 assembling the second arrays [are then assembled] into the EESU final assembly. Page 5

- 2. (Currently amended) [An] The method for making the electrical-energy-storage unit as recited in claim 1 wherein [the method of said claim provide] a second coating of glass is provided onto the double-coated composition-modified barium titanate powder being in contact with the nickel electrodes and having an applied working voltage of 3500 V across the parallel electrodes.
- 3. (Cancelled)
- 4. (Currently amended) [An] The method for making the electrical-energy-storage unit as recited in Claim 1 wherein [the method of said claim provide a unique double coating of the composition-modified barium titanate powder and the hot isostatic pressing at the near-minimum-temperature viscous-flow condition of the glass, 700° C,] a dielectric voltage breakdown strength of 5.0 x 10° V/cm was achieved across the electrodes of the components.
- 5. (Previously presented) [An] The method for making the electrical-energy-storage unit as recited in Claim 1 wherein the method [of said claim] provides an ease of manufacturing due to the softening temperature of the calcium magnesium aluminosilicate glass allowing the relatively low hot-isostatic-pressing temperatures of 700° C which in turn [provided] provides a void-free ceramic body.
- 6. (Previously presented) [An] The method for making the electrical-energy-storage unit as recited in Claim 1 wherein the method [of said claim] provides an ease of fabrication due to the softening temperature of the calcium magnesium aluminosilicate glass allowing the

Art Unit: 1775

relatively low hot-isostatic-pressing temperatures of 700° C which in turn allows the use of nickel for the conduction-path electrodes rather than expensive platinum, palladium, or palladium-silver alloy.

- 7. (Currently amended) [An] The method for making the electrical-energy-storage unit as recited in Claim 1 wherein the method [of said claim] provides an ease of fabrication due to the softening temperature of the calcium magnesium aluminosilicate glass allowing the relatively low hot-isostatic-pressing temperatures of 700°C, which feature along with the coating method provided a uniform-thickness shell of the calcium magnesium aluminosilicate glass and in turn provides hot-isostatic-pressed double-coated composition-modified barium titanate high-relative-permittivity layers that [were] are uniform and homogeneous in microstructure.
- 8. (Currently amended) [An] The method for making the electrical-energy-storage unit as recited in Claim 1 wherein the method [of said claim] provides the double coating of the basis particles of the composition-modified barium titanate powder [reduced] thereby reducing the leakage and aging of this material by an order of magnitude of the specification of this basis material, thus reducing the discharge rate to 0.1% per 30 days.
- 9. (Cancelled)
- 10. (Currently amended) [An] The method for making the electrical-energy-storage unit as recited in Claim 1 wherein the method [of said claim] provides a double coating of the composition-modified barium titanate powder, the hot-isostatic-pressing process, the high-density solder-bump packaging, and along with the double-layered array

Art Unit: 1775

configuration stored 52,220 W•h of electrical energy in a 2005 inches³ container.

- 11. (Currently amended) [An] The method for making the electrical-energy-storage unit as recited in Claim 1 wherein the method [of said claim] provides materials used: water-soluble precursors of barium (Ba), calcium (Ca), titanium (Ti), zirconium (Zr), manganese (Mn), yttrium (Y), neodymium (Nd), forming the composition-modified barium titanate powder, and the metals: nickel (Ni), and copper (Cu), which are not explosive, corrosive, or hazardous.
- 12. (Currently amended) [An] The method for making the electrical-energy-storage unit as recited in Claim 1 wherein the method [of said claim] provides an EESU that is not explosive, corrosive, or hazardous and therefore is a safe product when used in electrical vehicles, which include bicycles, tractors, buses, cars, or any device used for transportation or to perform work.
- 13. (Currently amended) [An] The method for making the electrical-energy-storage unit as recited in Claim 1 wherein the method [of said claim] provides an EESU which can store electrical energy generated from solar voltaic cells or other alternative sources for residential, commercial, or industrial applications.
- 14. (Currently amended) [An] The method for making the electrical-energy-storage unit as recited in Claim 1 wherein the method [of said claim] provides an[d] EESU which can store electrical energy from the present utility grid during the night when the demand for electrical power is low and then deliver the electrical energy during the peak power demand times and thus provide an effective power averaging function.

Art Unit: 1775

15. (Canceled)

16. (Canceled)

- 17. (Currently amended) [An] The method for making the electrical-energy-storage unit as recited in Claim 1 wherein the method [of said claim] provides a double coating of the composition-modified barium titanate powder and a hot-isostatic-pressing process which together [assisted] assists in allowing an applied voltage of 3500 V to a dielectric thickness of 12.76 x 10⁻⁶ m to be achieved.
- 18. (Currently Amended) [An] The method for making the electrical-energy-storage unit as recited in Claim 1 wherein the method [of said claim] provides a EESU which [was] when fully discharged and recharged, [and] the EESU's initial specifications [was] are not degraded.
- 19. (Currently Amended) [An] The method for making the electrical-energy-storage unit as recited in Claim 1 wherein the method [of said claim] provides a EESU which can be [was] safely charged to 3500 V and store[d] at least 52.22 kW•h of electrical energy.
- 20. (Currently Amended) [An] The method for making the electrical-energy-storage unit as recited in Claim 1 wherein the method [of said claim] provides a EESU that has a total capacitance of at least 31 F.
- 21. (Currently Amended) [An] The method for making the electrical-energy-storage unit as recited in Claim 1 wherein the method [of said claim] provides a EESU that [was] can be rapidly charged without damaging the material or reducing its life.

Allowable Subject Matter

- 2. Claims 1, 2, 4-8, 10-14 and 17-21 are allowed over the prior art of record.
- 3. The following is an examiner's statement of reasons for allowance:

The primary reason for allowance, is applicants' agreement to an examiner's amendment converting the product by process claims to method of making claims. While certain prior art of record, such as US Patent 6,268,054 (Costantino et al.), discloses a dielectric layers for multilayer ceramic capacitors wherein the dielectric material is made up oxide coated barium titanate particles and US Patent 6,078,494 (Hansen), which discloses a capacitor having a dielectric material made of a calcined powder of doped barium-calcium-zirconium titanate with nickel or nickel alloy electrodes, the prior art of record fails to teach or suggest the claimed steps for making the electrical energy storage unit.

Any comments considered necessary by applicant must be submitted no later than the payment of the issue fee and, to avoid processing delays, should preferably accompany the issue fee. Such submissions should be clearly labeled "Comments on Statement of Reasons for Allowance."

Conclusion

4. Any inquiry concerning this communication or earlier communications from the examiner should be directed to Stephen Stein whose telephone number is 572-272-1544. The examiner can normally be reached on Monday through Friday from 8:30 a.m. to 5:00 p.m. If the attempts to reach the examiner are unsuccessful, the examiner's supervisor, Deborah Jones can be reached by dialing 571-272-1535. The official fax number is 703-872-9306.

Art Unit: 1775

Page 10

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).

September 28, 2004

Stephen J. Stein Primary Examiner

Art Unit 1775